WHAT IS CLAIMED IS:

- 1. A stable, phase-pure magnesium-substituted crystalline hydroxyapatite comprising from about 2.0 to about 29 wt% magnesium, wherein at least 75 wt% of the magnesium content is substituted for calcium ions in the hydroxyapatite lattice structure.
- 5 2. The phase-pure magnesium-substituted crystalline hydroxyapatite of Claim 1, comprising from about 3.5 to about 28.4 wt% magnesium.
 - 3. The phase-pure magnesium-substituted crystalline hydroxyapatite of Claim 2, comprising from about 5 to about 25 wt% magnesium.
 - 4. The phase-pure magnesium-substituted crystalline hydroxyapatite of Claim 1, wherein essentially all of the magnesium content is substituted for calcium ions in the hydroxyapatite lattice structure.
 - 5. The phase-pure magnesium-substituted crystalline hydroxyapatite of Claim 1, comprising crystal agglomerates having a particle size between about 5 nm and about 100 microns.
- 15 6. A method for the preparation of phase-pure crystalline magnesium-substituted hydroxyapatite comprising mechanochemically reacting a source of calcium ions, a source of magnesium ions, a source of phosphate ions and a source of hydroxide ions, at least one of which is soluble in water, in a aqueous reaction medium until said magnesium substituted-hydroxyapatite is formed.
- 7. The method of Claim 6, wherein said ion sources are stoichiometrically selected to provide a predetermined level of magnesium substitution.

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- 8. The method of Claim 6, further comprising the step of separating said magnesium-substituted hydroxyapatite from said aqueous reaction medium.
- 9. The method of Claim 8, further comprising the step of washing said magnesiumsubstituted hydroxyapatite with water.
- The method of Claim 9, further comprising the step of drying said magnesium-substituted hydroxyapatite.
 - 11. The method of Claim 9, further comprising the step of washing said magnesium-substituted hydroxyapatite with an aqueous ammonium citrate solution before washing said magnesium- substituted hydroxyapatite with water.
 - 12. The method of Claim 6, wherein at least one of the ion sources is water-insoluble.
 - 13. The method of Claim 12, wherein the calcium ion source or the magnesium ion source is water-insoluble.
 - 14. The method of Claim 6, wherein said phosphate ion source is water soluble.
- 15. The method of Claim 6, wherein said magnesium ion source is selected from the group consisting of magnesium hydroxide, magnesium carbonate, magnesium halides, magnesium oxide, magnesium nitrate and magnesium phosphate.
 - 16. The method of Claim 15, wherein said magnesium ion source is magnesium hydroxide.
 - 17. The method of Claim 6, wherein said calcium ion source is selected from the group consisting of calcium hydroxide, calcium carbonate, calcium halides, calcium oxide, calcium nitrate and calcium phosphate.

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- 18. The method of Claim 17, wherein said calcium ion source is calcium hydroxide.
- 19. The method of Claim 6, wherein said phosphate ion source is selected from the group consisting of ammonium phosphates, calcium phosphates, magnesium phosphates, and sodium phosphates.
- The method of Claim 19, wherein said phosphate ion source is diammonium hydrogen phosphate.
 - 21. A packing material for use in a chromatography column or gas sensor or as a catalytic support comprising the magnesium-substituted hydroxyapatite of Claim 1.
 - 22. A biocompatible hard tissue implant comprising the magnesium-substituted hydroxyapatite of Claim 1.
 - 23. The biocompatible hard tissue implant of Claim 22, comprising a metal or polymeric implant coated with said magnesium-substituted hydroxyapatite.
 - 24. The biocompatible hard tissue implant of claim 22, comprising a polymeric composite.
- 25. A granular fill for direct incorporation into human or animal tissues comprising the magnesium-substituted hydroxyapatite of Claim 1.
 - 26. The granular fill of claim 25, comprising a metal or polymeric composite for filling dental cavities.
 - 27. A plant growth substrate comprising the magnesium-substituted hydroxyapatite of Claim 1.

A dentifrice composition comprising the magnesium-substituted hydroxyapatite of

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claim 1.

- 29. A method for increasing the magnesium content in the lattice structure of magnesium-substituted crystalline hydroxyapatite relative to the calcium content of the lattice structure and to the non-lattice magnesium content, said method comprising washing said magnesium-substituted hydroxyapatite with an aqueous ammonium citrate solution.
- 30. A host material for luminescent applications comprising the magnesium substituted hydroxyapatite of claim 1.